California Medical Device Recall Information

Recall Name
CareFusion 303 Recalls Alaris Pump Module Due To Possible Malfunction

Recall Date	Product Description	Recalling Firm	Recall Reason
6/15/12	Alaris Pump Module, Model 8100	**CareFusion 303, Inc.** San Diego, CA	*Suspected of keypad separating from door assembly leading to potential fluid ingress and malfunction*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Alaris Pump Module, Model 8100 Serial Numbers affected: Affected Serial Number List	**CA**, nationwide	Manufactured from October, 2011 through February, 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm316612.htm